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* * * * * Welcome to STN International * * * * *

NEWS	1		Web Page URLs for STN Seminar Schedule - N. America
NEWS	2	Apr 08	"Ask CAS" for self-help around the clock
NEWS	3	Apr 09	BEILSTEIN: Reload and Implementation of a New Subject Area
NEWS	4	Apr 09	ZDB will be removed from STN
NEWS	5	Apr 19	US Patent Applications available in IFICDB, IFIPAT, and
IFIUDB			
NEWS	6	Apr 22	Records from IP.com available in CAPLUS, HCAPLUS, and
ZCAPLUS			
NEWS	7	Apr 22	BIOSIS Gene Names now available in TOXCENTER
NEWS	8	Apr 22	Federal Research in Progress (FEDRIP) now available
NEWS	9	Jun 03	New e-mail delivery for search results now available
NEWS	10	Jun 10	MEDLINE Reload
NEWS	11	Jun 10	PCTFULL has been reloaded
NEWS	12	Jul 02	FOREGE no longer contains STANDARDS file segment
NEWS	13	Jul 22	USAN to be reloaded July 28, 2002; saved answer sets no longer valid
NEWS	14	Jul 29	Enhanced polymer searching in REGISTRY
NEWS	15	Jul 30	NETFIRST to be removed from STN
NEWS	16	Aug 08	CANCERLIT reload
NEWS	17	Aug 08	PHARMAMarketLetter(PHARMAML) - new on STN
NEWS	18	Aug 08	NTIS has been reloaded and enhanced
NEWS	19	Aug 19	Aquatic Toxicity Information Retrieval (AQUIRE) now available on STN
NEWS	20	Aug 19	IFIPAT, IFICDB, and IFIUDB have been reloaded
NEWS	21	Aug 19	The MEDLINE file segment of TOXCENTER has been reloaded
NEWS	22	Aug 26	Sequence searching in REGISTRY enhanced
NEWS	23	Sep 03	JAPIO has been reloaded and enhanced
NEWS	24	Sep 16	Experimental properties added to the REGISTRY file
NEWS	25	Sep 16	CA Section Thesaurus available in CAPLUS and CA
NEWS	26	Oct 01	CASREACT Enriched with Reactions from 1907 to 1985
NEWS	27	Oct 21	EVENTLINE has been reloaded
NEWS	28	Oct 24	BEILSTEIN adds new search fields
NEWS	29	Oct 24	Nutraceuticals International (NUTRACEUT) now available on
STN			
NEWS	30	Oct 25	MEDLINE SDI run of October 8, 2002
NEWS	31	Nov 18	DKILIT has been renamed APOLLIT
NEWS	32	Nov 25	More calculated properties added to REGISTRY
NEWS	33	Dec 02	TIBKAT will be removed from STN
NEWS	34	Dec 04	CSA files on STN
NEWS	35	Dec 17	PCTFULL now covers WP/PCT Applications from 1978 to date
NEWS	36	Dec 17	TOXCENTER enhanced with additional content
NEWS	37	Dec 17	Adis Clinical Trials Insight now available on STN
NEWS	38	Dec 30	ISMEC no longer available

NEWS 39 Jan 13 Indexing added to some pre-1967 records in CA/CAPLUS
 NEWS 40 Jan 21 NUTRACEUT offering one free connect hour in February 2003
 NEWS 41 Jan 21 PHARMAML offering one free connect hour in February 2003
 NEWS 42 Jan 29 Simultaneous left and right truncation added to COMPENDEX,
 ENERGY, INSPEC
 NEWS 43 Feb 13 CANCERLIT is no longer being updated
 NEWS 44 Feb 24 METADEX enhancements
 NEWS 45 Feb 24 PCTGEN now available on STN
 NEWS 46 Feb 24 TEMA now available on STN
 NEWS 47 Feb 26 NTIS now allows simultaneous left and right truncation
 NEWS 48 Feb 26 PCTFULL now contains images
 NEWS 49 Mar 04 SDI PACKAGE for monthly delivery of multifile SDI results

NEWS EXPRESS January 6 CURRENT WINDOWS VERSION IS V6.01a,
 CURRENT MACINTOSH VERSION IS V6.0b(ENG) AND V6.0Jb(JP),
 AND CURRENT DISCOVER FILE IS DATED 01 OCTOBER 2002

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* * * * * STN Columbus * * * * *

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=> file caplus

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FILE 'CAPLUS' ENTERED AT 10:11:23 ON 13 MAR 2003

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FILE COVERS 1907 - 13 Mar 2003 VOL 138 ISS 11
FILE LAST UPDATED: 12 Mar 2003 (20030312/ED)

This file contains CAS Registry Numbers for easy and accurate substance identification.

```
=> s 1,3-cyclic glycerophosphate
      7465882 1
      5712254 3
      258238 CYCLIC
      8095 GLYCEROPHOSPHATE
L1      2 1,3-CYCLIC GLYCEROPHOSPHATE
      (1(W)3(W)CYCLIC(W)GLYCEROPHOSPHATE)
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=> d 11 1-2 all
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L1 ANSWER 1 OF 2 CAPLUS COPYRIGHT 2003 ACS
AN 2000:706969 CAPLUS
DN 133:261536
TI Pharmaceutical compositions comprising cyclic glycerophosphates and
   analogs thereof for promoting neural cell differentiation
IN Shinitzky, Meir
PA Yeda Research and Development Co. Ltd., Israel
SO PCT Int. Appl., 42 pp.
   CODEN: PIXXD2
DT Patent
LA English
IC ICM A61K031-00
CC 1-11 (Pharmacology)
   Section cross-reference(s): 29, 63
```

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2000057865	A2	20001005	WO 2000-IL185	20000324
	WO 2000057865	A3	20010628		
	W:	AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CR, CU, CZ, DE, DK, DM, DZ, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM			
	RW:	GH, GM, KE, LS, MW, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG			
	BR 2000009296	A	20011218	BR 2000-9296	20000324
	EP 1162959	A2	20011219	EP 2000-912877	20000324
	R:	AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO			
	JP 2002540146	T2	20021126	JP 2000-607616	20000324
PRAI	IL 1999-129178	A	19990325		
	WO 2000-IL185	W	20000324		
OS	MARPAT 133:261536				

- AB Cyclic glycerophosphates and analogs thereof (CGs) are shown to exert neural promoting activities in target cells. Such activities include promotion of neuronal outgrowth, promotion of nerve growth, provision of dopaminotrophic supporting environment in a diseased portion of the brain, prevention of nerve degeneration and nerve rescue. These activities of the CGs render them useful for treatment of various disorders including but not limited to mental disorders such as, for example, schizophrenia, dementia or disorders resulting in learning disabilities. In addn., these CGs may be used for the treatment of neurodegenerative conditions such as Alzheimer's disease, Parkinson's disease, conditions resulting from exposure to harmful environmental factors or resulting from a mech. injury. The CGs may also be used to treat an individual suffering from a primary neurodegenerative condition in order to prevent or reduce the appearance of secondary degeneration in addnl. nerves ("nerve rescue"). For example, neural outgrowth of PC12 cells was seen when cells were grown in the presence of nerve growth factor (50 ng/mL) or 1,3-cyclic glycerophosphate (1 .mu.M), but not in the presence of linear .alpha.-glycerophosphate.
- ST cyclic glycerophosphate neuronal differentiation mental disorder; antipsychotic schizophrenia cyclic glycerophosphate; Alzheimer disease parkinsonism cyclic glycerophosphate
- IT Anti-Alzheimer's agents
Antiparkinsonian agents
Antipsychotics
Mental disorder
Nervous system agents
Schizophrenia
(compns. comprising cyclic glycerophosphates for promoting neural differentiation for therapeutic uses)
- IT Monoamines
Neurotrophic factors
RL: BPR (Biological process); BSU (Biological study, unclassified); BIOL (Biological study); PROC (Process)
(compns. comprising cyclic glycerophosphates for promoting neural differentiation for therapeutic uses)
- IT Nerve
(degeneration, prevention of; compns. comprising cyclic glycerophosphates for promoting neural differentiation for therapeutic uses)
- IT Mental disorder
(dementia; compns. comprising cyclic glycerophosphates for promoting neural differentiation for therapeutic uses)
- IT Nerve
(differentiation; compns. comprising cyclic glycerophosphates for promoting neural differentiation for therapeutic uses)
- IT Learning
(disorder; compns. comprising cyclic glycerophosphates for promoting neural differentiation for therapeutic uses)
- IT Nerve
(dopaminergic, degeneration of; compns. comprising cyclic glycerophosphates for promoting neural differentiation for therapeutic uses)
- IT Cell differentiation

(inducers; compns. comprising cyclic glycerophosphates for promoting neural differentiation for therapeutic uses)

IT Nerve, disease
(injury, neuronal rescue after; compns. comprising cyclic glycerophosphates for promoting neural differentiation for therapeutic uses)

IT Cell differentiation
Cell differentiation
(neuronal; compns. comprising cyclic glycerophosphates for promoting neural differentiation for therapeutic uses)

IT Drug delivery systems
(oral; compns. comprising cyclic glycerophosphates for promoting neural differentiation for therapeutic uses)

IT Drug delivery systems
(osmotic pumps; compns. comprising cyclic glycerophosphates for promoting neural differentiation for therapeutic uses)

IT Cell proliferation
(promotion of; compns. comprising cyclic glycerophosphates for promoting neural differentiation for therapeutic uses)

IT Drug delivery systems
(topical; compns. comprising cyclic glycerophosphates for promoting neural differentiation for therapeutic uses)

IT 298701-05-0P
RL: BAC (Biological activity or effector, except adverse); BPN (Biosynthetic preparation); BSU (Biological study, unclassified); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)
(compns. comprising cyclic glycerophosphates for promoting neural differentiation for therapeutic uses)

IT 711-07-9P 13507-10-3P 22227-09-4P 118897-32-8P 123406-35-9P 286020-33-5P 298701-06-1P 298701-08-3P 298701-09-4P 298701-78-7P
RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)
(compns. comprising cyclic glycerophosphates for promoting neural differentiation for therapeutic uses)

IT 51-61-6, Dopamine, biological studies 59-92-7, biological studies 102-32-9, DOPAC 306-08-1, Homovanillic acid
RL: BPR (Biological process); BSU (Biological study, unclassified); BIOL (Biological study); PROC (Process)
(compns. comprising cyclic glycerophosphates for promoting neural differentiation for therapeutic uses)

IT 9001-86-9, Phospholipase C
RL: BUU (Biological use, unclassified); BIOL (Biological study); USES (Uses)
(compns. comprising cyclic glycerophosphates for promoting neural differentiation for therapeutic uses)

IT 57-55-6, 1,2-Propanediol, reactions 96-26-4, Dihydroxyacetone 504-63-2, 1,3-Propanediol 770-12-7, Phenyl phosphorodichloridate 819-83-0, Disodium .beta.-glycerophosphate 4799-67-1 14690-00-7, 2-Benzoyloxy-1,3-propanediol 22002-87-5 26776-70-5, Dihydroxyacetone dimer
RL: RCT (Reactant); RACT (Reactant or reagent)
(compns. comprising cyclic glycerophosphates for promoting neural differentiation for therapeutic uses)

differentiation for therapeutic uses)

IT 187976-16-5P
 RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)
 (compns. comprising cyclic glycerophosphates for promoting neural differentiation for therapeutic uses)

L1 ANSWER 2 OF 2 CAPLUS COPYRIGHT 2003 ACS
 AN 1993:534139 CAPLUS
 DN 119:134139
 TI Formation of 1,3-cyclic glycerophosphate by the action of phospholipase C on phosphatidylglycerol
 AU Shinitzky, Meir; Friedman, Peter; Haimovitz, Rachel
 CS Dep. Membrane Res. Biophys., Weizmann Inst. Sci, Rehovot, 76100, Israel
 SO Journal of Biological Chemistry (1993), 268(19), 14109-15
 CODEN: JBCHA3; ISSN: 0021-9258
 DT Journal
 LA English
 CC 7-3 (Enzymes)
 AB The action of phospholipase C (PLC) from *Bacillus cereus* on phosphatidylglycerol (PG), derived from egg yolk phosphatidylcholine (PC), was examd. in an ether-water mixt. The PLC cleavage of PG and PC followed a Michaelis-Menten kinetics with apparent Vmax values per 1 .mu.g enzyme of 0.26 and 0.91 .mu.mol.min⁻¹ and Km values of 10 and 12 mM, resp. When the same enzymic reaction was carried out in minimally buffered aq. soln. of 1% Triton X-100, the decrease in pH with respect to phospholipid cleavage was as expected with PC but much less pronounced with PG. This could be accounted for by .alpha.-glycerophosphate, in the PLC hydrolysis of PG. Examn. of the chem. nature of the water-sol. product of PG by 31P NMR revealed a single band at 2.31 ppm, while the bands of .alpha.-glycerophosphate and .beta.-glycerophosphate appeared at 5.12 and 4.57 ppm, resp. Basic hydrolysis of the phospholipase cleavage product of PG (0.1 M NaOH for 1 min at 80 .degree.C) followed by neutralization shifted its 31P NMR band to 5.18 ppm, which practically coincided with that of .alpha.-glycerophosphate. Analogous expts. were carried out with PG labeled with 3H at the carbon 2 of the glycerol headgroup ([3H]PG). Autoradiog. of thin layer chromatog. (TLC) of the [3H]PG enzymic hydrolyzate displayed a single 3H-labeled compd., which could be converted to .alpha.-glycerophosphate by basic hydrolysis. These results strongly suggest that the phosphate headgroup of PG is cleaved off by PLC as 1,3-cyclic glycerophosphate. A series of PLC expts. with phosphatidyl dihydroxyacetone and phosphatidyl 1,3-propanediol as model substrates supported this assignment. Two-dimensional homonuclear 1H NMR correlated spectra as well as IR spectra carried out on the isolated sodium salt of this product could further confirm such a structure. The unique structure and chem. nature of 1,3-cyclic glycerophosphate may bear a distinct physiol. function.

ST cyclic glycerophosphate formation phospholipase C phosphatidylglycerol
 IT Phosphatidylglycerols
 RL: RCT (Reactant); RACT (Reactant or reagent)

(cleavage of, by phospholipase C, cyclic glycerophosphate formation
in)
IT Michaelis constant
(of phospholipase C, with phosphatidylglycerol)
IT Phosphatidylcholines, reactions
RL: RCT (Reactant); RACT (Reactant or reagent)
(reaction of, with phospholipase C, kinetics of, phosphatidylglycerol
in relation to)
IT 9001-86-9, Phospholipase C
RL: BIOL (Biological study)
(cyclic glycerophosphate formation by, of Bacillus cereus, in
phosphatidylglycerol cleavage)
IT 42320-97-8
RL: FORM (Formation, nonpreparative)
(formation of, by phospholipase C cleavage of phosphatidylglycerol)
IT 149864-37-9
RL: FORM (Formation, nonpreparative)
(formation of, by phospholipase C cleavage of
phosphatidylhydroxyacetone)
IT 13507-10-3
RL: FORM (Formation, nonpreparative)
(formation of, by phospholipase C cleavage of phosphatidylpropanediol)

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